CLAIMS

1. A pharmaceutical composition comprising a solid intimate mixture of human growth releasing factor (GRF) and a stabilizing amount of saccharose, alone or in combination with other excipients.

2. The pharmaceutical composition according to Claim 1, wherein the solid intimate mixture is a lyophilizate.

3. The pharmaceutical composition according to any of Claims 1 to 2, wherein the stabilizing agent is saccharose alone.

4. The pharmaceutical composition according to any of claims 1 to 3, containing 3 or 10 mg/vial of hGRF.

5. The pharmaceutical composition according to any of Claims 1 to 4 comprising 3 or 10 mg/vial of hGRF and 20.52 or 68.4 mg/vial of saccharose.

6. The pharmaceutical composition according to any of Claims 1 to 5 further comprising buffering agents.

7. A process for preparing a pharmaceutical composition according to any of Claims 1 to 6, comprising the preparation of an aqueous solution of the components, the distribution within containers and the lyophilization in the containers.

8. Forms of presentation of said pharmaceutical composition comprising the solid mixture according to any of Claims 1 to 6, hermetically closed in a sterile condition within a container suited for storage before use and for reconstitution of the mixture into a solvent or into a solution for injectables.

9. A solution comprising the solid mixture according to any of Claims 1 to 6, reconstituted in a solvent or a solution for injectables.

10. The solution according to any of Claim 9, wherein the pH in comprised between 4 and 6.